## Invited Talk

at

## International Symposium on Electron-Molecule Collisions and Swarms

Tokyo, Japan

18-20 July, 1999

## High Resolution UV Emission Spectroscopy of Molecules Excited by Electron Impact

G. K. James, J. M. Ajello, L. Beegle, M. Ciocca<sup>w</sup>, D. Dziczek, I. Kanik, C. Noren, C. Jonin and D. Hansen

Jet Propulsion Laboratory, California Institute of Technology, Pasadena, CA 91109, USA \*\*Department of Physics and Astronomy, University of Kentucky, Lexington KY 40506, USA

Photodissociation via discrete line absorption into predissociating Rydberg and valence states is the dominant destruction mechanism of CO and other molecules in the interstellar medium and molecular clouds. Accurate values for the rovibronic oscillator strengths of these transitions and predissociation yields of the excited states are required for input into the photochemical models that attempt to reproduce observed abundances. We report here on our latest experimental results of the electron collisional properties of CO and N<sub>2</sub> obtained using the 3-meter high resolution single-scattering spectroscopic facility at JPL.

The Fourth Positive (A  ${}^{1}\Pi - X {}^{1}\Sigma^{+}$ ) band system of CO is ubiquitously observed in astronomy, appearing in UV spectra from the sun, the planets, and the interstellar medium. We have measured laboratory emission spectra of CO at medium resolution (~0.03 nm full width at half maximum (FWHM)) produced by electron impact excitation at 20 eV and 100 eV in the wavelength range 130 to 205 nm. Features observed in these far-ultraviolet fluorescence spectra correspond to the Fourth Positive band system, atomic multiplets from C and O, together with their ions. The spectral region between 130 and 180 nm observed at 100 eV is shown in Figure 1. Electron impact emission cross sections measured in the present work (and listed in Beegle et al., 1998) are about 10% higher than the previous measurements of Aarts and de Heer (1970), Mumma et al. (1971), and Ajello (1971). For example, the cross section at 100 eV for the entire Fourth Positive band system as reported by Ajello (1971) was ~31 x  $10^{-18}$  cm², compared to a result of 34.4 x  $10^{-18}$  cm² obtained in the present work. However, the value of the studies from 25 years ago which were in close agreement has been undermined by the different calibration standards employed, all of which have undergone substantial revision. In addition, the contribution of cascade to the emission cross section was underestimated in the earlier works.

The present CO Fourth Positive spectral data were used in combination with those of DeLeon (1989) to determine the dependence of the electronic transition moment ( $R_e$ ) on the internuclear distance ( $r_{v',v''}$ ). The transition moment data of DeLeon (1989) were normalized to our values of  $R_e$  in the 1.35-1.40 Å region, as shown in Figure 2. The combined data set was then fitted with a polynomial out to an internuclear distance of 1.8 Å. This curve has the form:

$$R_e = 7.64 (1-0.957 r_{v'v''} + 0.2247 r^2_{v'v''}),$$

and is in good agreement with recent work of others (Federman et al. (1997), Smith et al. (1994), Chan et al. (1993), and Kirby and Cooper (1989)). This agreement supports the conclusions of Morton and Noreau (1994) who recommended the use of the CO Fourth Positive band oscillator strengths measured by Chan et al. (1993).

١

level of the  $N_2$  c' state will change with temperature, from very small in the Earth's atmosphere (where temperatures approach 1000 °K (*Hedin*, 1987)) to much larger in the atmospheres of Triton and Titan (where temperatures are about 50-200 °K). The opposite temperature trend will occur for the v'=3 level. The results seem to establish a general trend of increasing mean predissociation yield with vibrational level by including in the study the mean 300 °K predissociation yield of 15% from our work on the c' v'=0 level (*Shemansky et al.*, 1995). The other strong levels, v'= 2 and 6, also need to be studied at high spectral resolution.

The excitation cross section at 100 eV can be calculated based on the emission cross section and the mean predissociation yield at 175 °K (42% and 50 % for v'= 3 and 4, respectively). The resulting total excitation cross section values are 13.6 x 10<sup>-19</sup> cm² for v' = 3, and 19.8 x 10<sup>-19</sup> cm² for v' = 4. Although these absolute cross sections are dependent on normalization to the predissociation yield for J' =9 measured by Walter et al. (1994), the spectra obtained in this study are the first electron impact spectra to separate the effects of blending of the b' (v' = 10 and 13) levels with the c' (v' = 3 and 4) levels, respectively. The ratio of excitation cross sections c'(4)/c'(3) obtained here is 1.5. The corresponding excitation cross section ratio measured by Geiger and Schroder (1969) is 2.7, by Chan et al. (1993) is 2.6, by Zipf and McClaughlin (1978) is 2.9, and by Carter (1972) is 1.7. The last ratio was measured by high resolution absorption spectroscopy and the sets of former works were obtained by energy loss spectroscopy. The previously published cross section ratios from energy loss results must be viewed with caution, since the spectra did not resolve the structure from overlapping Rydberg and valence states.

## References:

Aarts, J. F. M. and de Heer, F. J., J Chem Phys, 52, 3554, 1970.

Ajello, J. M., J Chem Phys, 55, 3158, 1971.

Beegle, L., Ajello, J., James, G., Alvarez, M. and Dziczek, D., submitted to Astron. Astrophys, 1998. Carter, V., J. Chem. Phys., 56, 4195, 1972.

Chan, F., Cooper, G. and Brion, C. E., Chem Phys, 170, 123, 1993.

Chan, W. F., Cooper, G., Sodhi, R. and Brion, C., Chem. Phys., 170, 81, 1993.

De Leon, R. L., J. Chem. Phys., 91, 5859, 1989.

Federman, S. R., Menningen, K. L., Lee, W. and Stoll, J. B., ApJ, 477, L61, 1997.

Geiger, J. and Schroder, B., J. Chem. Phys., 50, 7, 1969.

Hedin, A. E., J. Geophys. Res., 92, 4649, 1987.

Kanik, I., James, G.K. and Ajello, J.M., Phys. Rev. A, 51, 2067, 1995.

Kirby, K. and Cooper, D.L., J. Chem. Phys., 90, 4895, 1989.

Morton, D. C. and Noreau, L., Ap.J.Supp., 95, 301, 1994.

Mumma, M. J., Stone, E. J. and Zipf, E. C., J. Chem. Phys., 54, 2627, 1971.

Shemansky, D. E., Kanik, I. and Ajello, J. M., Ap. J., 452, 480, 1995.

Smith, P. L., Stark, G., Yoshino, K. and Ito, K., Ap. J., 431, L413, 1994.

Walter, C. W., Cosby, P. C. and Helm, H., Phys. Rev. A, 50, 2930, 1994

Zipf, E. C. and McLaughlin, R.W., Planet. Space Sci., 26, 449, 1978.

Acknowledgments: This work was supported by NASA, NSF, and AFOSR grants.